

meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at <http://www.fda.gov/AdvisoryCommittees/Calendar/default.htm>. Scroll down to the appropriate advisory committee link.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before November 23, 2011. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before November 15, 2011. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by November 16, 2011.

Persons attending FDA's advisory committee meetings are advised that the Agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Kalyani Bhatt at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at <http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm> for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: September 20, 2011.

Jill Hartzler Warner,

Acting Associate Commissioner for Special Medical Programs.

[FR Doc. 2011-24532 Filed 9-22-11; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2010-N-0318]

Novartis Pharmaceuticals Corp. et al.; Withdrawal of Approval of 27 New Drug Applications and 58 Abbreviated New Drug Applications; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of July 21, 2010 (75 FR 42455). The document withdrew approval of 27 new drug applications (NDAs) and 58 abbreviated new drug applications (ANDAs) from multiple applicants. The published document excluded a footnote in the table. This document corrects that error.

FOR FURTHER INFORMATION CONTACT:

Joyce Strong, Office of Policy, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, rm. 3208, Silver Spring, MD 20993-0002, 301-796-9148.

SUPPLEMENTARY INFORMATION: In FR Doc. 2010-17785, appearing on page 42455, in the **Federal Register** of Wednesday, July 21, 2010, the following correction is made:

1. On page 42456, in Table 1, under the "Drug" column, correct the entry for "Proventil (albuterol USP) Inhalation Aerosol" to read "Proventil (albuterol USP) Inhalation Aerosol 1".

2. On page 42456, at the end of the table, add footnote number 1 to read:

This product included an oral pressurized metered-dose inhaler that contained chlorofluorocarbons (CFCs) as a propellant. CFCs may no longer be used as a propellant for any albuterol metered-dose inhalers. (See 70 FR 17168, April 4, 2005.)

Dated: September 19, 2011.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2011-24400 Filed 9-22-11; 8:45 am]

BILLING CODE 4160-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

National Practitioner Data Bank; Name Change of Proactive Disclosure Service (PDS) to Continuous Query

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice.

SUMMARY: On March 7, 2007, the Health Resources and Services Administration (HRSA), Department of Health and Human Services (HHS), published in the **Federal Register** a notice announcing the implementation of a prototype for querying the National Practitioner Data Bank (NPDB), then known as Proactive Disclosure Service (PDS). This notice announces that the prototype status is removed and that PDS is now known as Continuous Query.

DATES: The effective date of this status upgrade and name change is September 23, 2011.

FOR FURTHER INFORMATION CONTACT:

Director, Division of Practitioner Data Banks, Bureau of Health Professions, Health Resources and Services Administration, Parklawn Building, 5600 Fishers Lane, Room 8-103, Rockville, MD 20857; telephone number: (301) 443-2300.

SUPPLEMENTARY INFORMATION:

I. Background

On March 7, 2007, the National Practitioner Data Bank (NPDB) published in the **Federal Register** (72 FR 10227) a notice announcing a Proactive Disclosure Service (PDS) prototype. The PDS was offered as an alternative to the traditional querying of the NPDB and allowed for on-going monitoring of a practitioner's credentials. PDS is a subscription service that notifies subscribers, which are registered entities that are eligible to query the NPDB or the Healthcare Integrity and Protection Data Bank (HIPDB), of new information on any of their enrolled practitioners within 24 hours of the NPDB or HIPDB receipt of the information. The PDS prototype was available for enrollment beginning on April 30, 2007 to a select group of NPDB registered entities. A few months later PDS was opened to all NPDB registered entities, as well as to those registered in the HIPDB. In the last year (July 1, 2010 through June 30, 2011), 1,965 entities had practitioner enrollments through PDS versus 14,370 entities that submitted traditional queries on